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# Quantitative Investigations about the Surface Texture Characteristics of the Nickel-Carbon Composite Thin Films using Stereometric Analysis

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**Abstract**—The purpose of this study was to describe the 3-D nano-scaled surface texture of the Nickel-Carbon (Ni-C) nanocomposite thin films composed of Ni nanoparticles with different average sizes embedded in amorphous hydrogenated carbon using stereometric analysis. The 3-D nanoscaled surface texture was studied by atomic force microscopy (AFM) and stereometric analysis. In a wider context, we also discuss our findings with regard to manufacture of the Nickel-Carbon composite thin films with desired surface characteristics.

**Keywords**—atomic force microscopy; Nickel-Carbon composite thin films; stereometric analysis; surface micromorphology

## I. INTRODUCTION

Nickel is conductor, hard and ferromagnetic at room temperature. Due to a layer of its oxide acting as a protection, it is corrosion-resistant, so its thin layers are used to cover other metals to prevent corrosion, which has many uses for the production of industrial tools [1-3].

Carbon nanocomposites containing metal nanoparticles in the carbon amorphous environment have attracted the attention of researchers owing to their interesting physical applications to investigate their application in various industrial and medical fields [4-6], and they can also be used as coating materials in electronics, spintronics, optics, biomedicine, and ferroelectric memories [7-9].

Thus, the researchers, with changes in the method and conditions for the construction of these thin layers or even small changes in the composite structure, create various physical properties in them. Therefore, control over the method, fabrication conditions, and impurities of metal-carbon thin film can lead to control over structural properties and, consequently, other physical properties [10-19].

A nickel-carbon thin film is considered and studied used as an optical coating, a catalyst, and a water splitting electrode [20, 21].

In this article, stereometric analyses have been carried out to investigate the 3-D surface texture of engineered composite

surfaces of Nickel-Carbon, which have been imaged using AFM.

## II. MATERIALS AND METHODS

### A. Materials

Ni-C composite thin films were prepared by the methodology described elsewhere [1]. Producing the nickel nanoparticles in carbon bed is performed using RF-PECVD in acetylene gas environment. In this system, the chamber is vacuumed using rotary and diffusion pumps up to  $10^{-5}$  bar, and the chamber consists of two electrodes, one as the target electrode (nickel) and the other as a substrate. Next, the glass substrate with  $1 \times 1$  dimensions is placed inside the chamber on the electrode attached to earth, and acetylene gas enters the compartment to adjust the functioning pressure. The RF power is then applied to create the plasma and begin sputtering. For all samples (target: Nickel) were used next sputtering parameters: basic pressure  $10^{-5}$  [mbar]; work pressure 0.025 [mbar]; power 250 [W]. The deposition time was varied in different experiments for samples: A) 7 min; B) 10 min and C) 13 min.

### B. Methods

The parameters related to the surface texture of the films were investigated using atomic force microscopy (AFM) analysis. MountainsMap® Premium software was used in the analysis of surface topography [22]. The software makes use of advanced tools and is compatible with the AFM microscopy.

## III. RESULTS

The measurement data of the samples of three different deposition periods were entered into MountainsMap® Premium in order to analyze the samples (Figures: I, II and III).

First of all, the images were leveled by subtraction using the following method: the LS plane and shape deletion operation was employed.

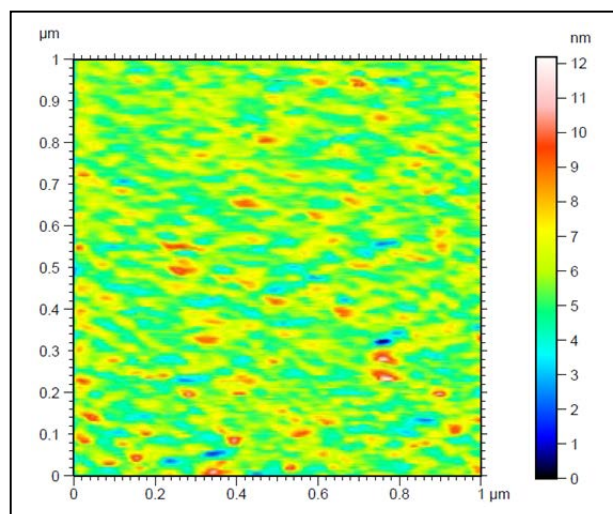


FIGURE I. THE AFM IMAGE OF THE SAMPLE: A

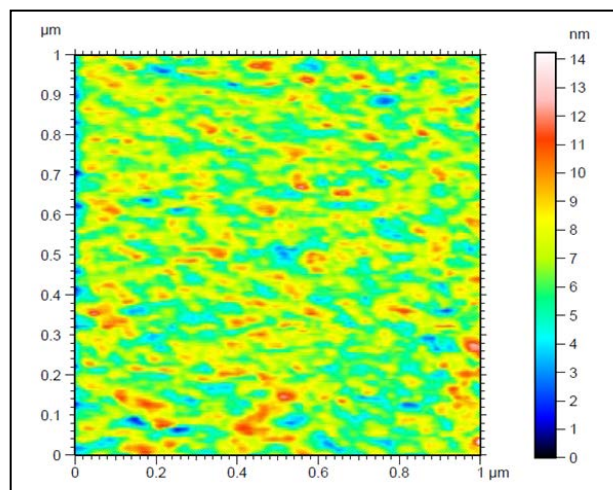


FIGURE II. THE AFM IMAGE OF THE SAMPLE: B

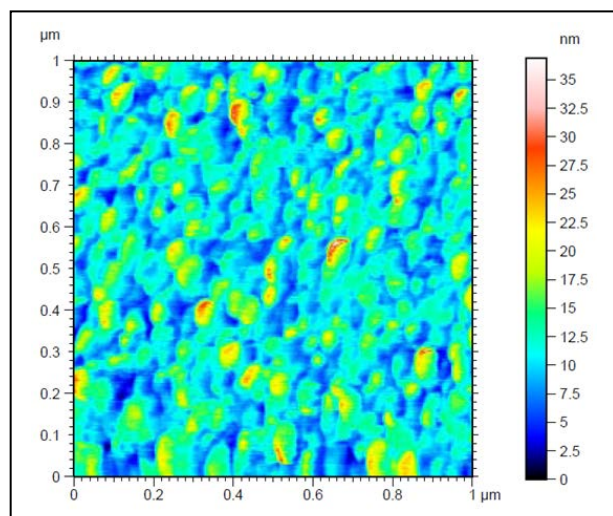


FIGURE III. THE AFM IMAGE OF THE SAMPLE: C

Secondly, identification cards including basic information about the samples (e.g. length, size, spacing, offset) were prepared. The data included in the cards were collected in Table I.

TABLE I. INFORMATION FROM THE IDENTIFICATION CARDS OF ALL THREE SAMPLES

	Sample 1	Sample 2	Sample 3
AXIS: X			
Length [μm]	1,00	1,00	1,00
Size [points]	256	256	256
Spacing [nm]	3,92	3,92	3,92
Offset [μm]	0,00	0,00	0,024
AXIS: Y			
Length [μm]	1,00	1,00	1,00
Size [lines]	256	256	256
Spacing [nm]	3,92	3,92	3,92
Offset [μm]	-1,0	-1,0	-1,02
AXIS: Z			
Length [nm]	12,2	14,2	0,00
Z min [nm]	-5,71	-6,8	0,00
Z max [nm]	6,48	7,44	0,00
Size [decimal place values]	159160	905994	0
Spacing [pm]	0,0766	0,0157	0,273

Thirdly, the 3-D image for each sample was generated. The image has a scale, graduation and axis system (Figures IV-VI).

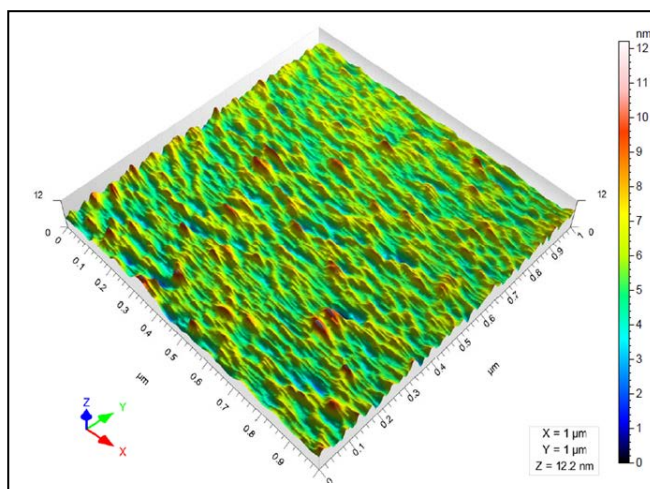


FIGURE IV. THE AFM 3D IMAGE OF SAMPLE A



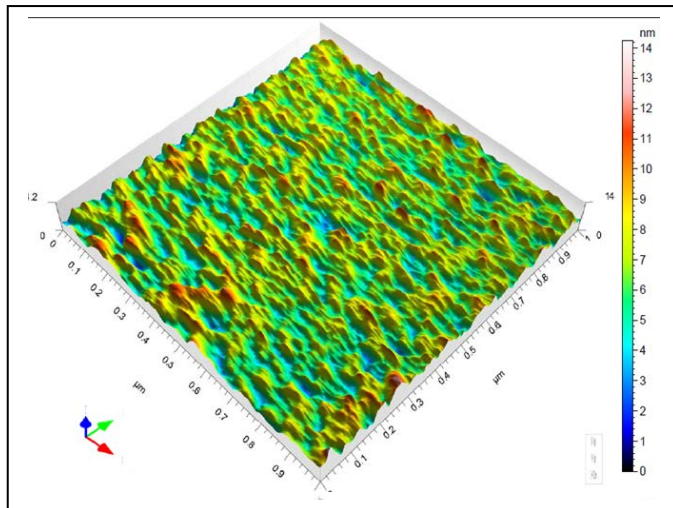


FIGURE V. THE AFM 3D IMAGE OF SAMPLE B

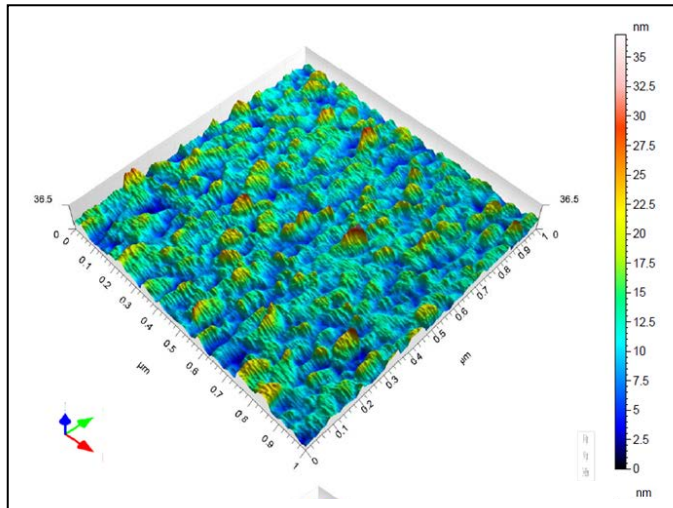


FIGURE VI. THE AFM 3D IMAGE OF SAMPLE C

After analyzing the surface samples, the values of surface stereometric parameters were collected (Table II). These parameters were determined according to ISO 25178 standard and EUR 15178N standard [23-30].

TABLE II. 3-D SURFACE STEREO-METRIC PARAMETERS

3-D surface stereometric parameters			
ISO 25178			
	Sample 1	Sample 2	Sample 3
<b>Sq</b> [nm]	0.999	1.43	3.76
<b>Ssk</b> [-]	0.524	0.102	0.795
<b>Sku</b> [-]	4.68	3.24	4.21
<b>Sp</b> [nm]	6.48	7.44	25.3
<b>Sv</b> [nm]	5.71	6.80	11.6
<b>Sz</b> [nm]	12.2	14.2	36.9
<b>Sa</b> [nm]	0.760	1.13	2.93
EUR 15178N			
<b>Sz</b> [nm]	9.68	10.3	31.5

#### IV. DISCUSSIONS

The deposition time for the samples was as the following: sample A: 7 minutes, sample B: 10 minutes and sample C: 13 minutes. After leveling and shape disposal, the images did not change, because the operations had been performed earlier.

The data from the identification cards for each sample are the same against X- and Y-axes. Only the offset value for the sample 3 is different. On the other hand, this information is more varied against the Z-axis. All the samples were logged in 256 points which fall 256 scanning lines. The measurements of each sample in X and Y-axes account 1  $\mu\text{m}$  x 1  $\mu\text{m}$ , and in Z-axis: 12.2 nm – sample A, 14.2 nm – sample B, 0.00 nm – sample C.

There were eight altitude parameters for the above samples, Table II).

The largest root mean height (Sq) shows the sample C (3.76 nm), whereas the lowest one has the sample A (0.999 nm). This parameter indicates the root mean square of a measured surface.

The surface skewness (Ssk) of all three samples is positive ( $Ssk > 0$ ). It indicates the dominance of peaks on their surface, and the highest number of them has the sample C (0.795), while the lowest number of peaks has the sample B (0.102).

The surface kurtosis (Sku) for each sample reaches the value above 3.00. Hence, it can be concluded that there are excessively high peaks or valleys on the surface with the highest value for the sample A (4.68). However, the lowest value appears in the sample B (3.24).

As far as the Sp and Sv parameters are considered, the highest maximum peak height and the largest pit height can be observed in the sample C ( $Sp = 25.3$  nm,  $Sv = 11.6$  nm), whereas the lowest values of these parameters are reached by the sample A ( $Sp = 6.48$  nm,  $Sv = 5.71$  nm). In addition, the sample C has the following values for the parameters:  $Sp = 7.44$  nm,  $Sv = 6.80$  nm.

On the basis of this information, it may be said that the greatest value of Sz parameter (i.e. the sum of the maximum peak height value and the maximum pit height value) has the sample C (36.9 nm), and the least value is observed in the sample A (12.2 nm).

The highest arithmetic mean surface height (Sa) also appears in the sample C (2.93 nm), while the lowest one in the sample A (0.760 nm). These values also represent the arithmetic means of vertical deviations of the surface from the mean surface.

The last obtained parameter was the Sz parameter (defined in the EUR 15178N standard) which was used to determine the heights of ten surface points. The sample C for this parameter shows the highest value (31.5 nm), and the sample A the lower value (9.68 nm). The sample with the deposition time of 13 minutes is found to be the sample that has the highest values of altitude parameters (except for the parameter Sku).

## V. CONCLUSION

The 3-D analysis of surface topography plays a significant role in the characteristics of material properties and functionality which may have various applications in technique. The stereometric analysis was conducted by means of MountainsMap® Premium software of the AFM images. It can be seen that the time deposition had an influence on the obtained values of the altitude parameters. The differences in the surface topography have been observed in each examined sample. The combination of RF-sputtering and RF-PECVD techniques represents a suitable route to prepare the Nickel–Carbon nanocomposite thin films composed of Ni nanoparticles with different average sizes embedded in amorphous hydrogenated carbon. Furthermore, all these morphological parameters can be used in various mathematical models to describe the local topography of the 3-D sample surface textures.

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